

ate member of the Broken Jug Formation (Fig. 2; Lucas and Lawton, 2000, fig. 3).

Bivalves

At NMMNH locality 4487, a coquina bed yields many bivalves catalogued as NMMNH P-32956–32961. These bivalves (Fig. 3A–H) are characterized by a small left valve that has a thick shell wall, is triangular in outline, and is clearly much larger than the right valve. The left valve is moderately convex to deeply excavated and is strongly arched dorsally. The beak is bluntly hooked, and the presence of a sulcus from the beak downward to produce a wing-like flare of the posteroventral margin is difficult to determine on most specimens due to poor preservation; however, it is visible on some specimens (e.g., Fig. 3D–E, H). The incurved beak is prominent but incomplete on all specimens due to abrasion. Those specimens with preserved external shell morphology show acute, imbricate, concentric growth lines and undulations (Fig. 3D). The shell is higher than long; left valve heights range from 30 to 35 mm (1.2–1.4 inches), and left valve lengths are much more variable, ranging from 10 to 30 mm (0.4–1.4 inches).

These specimens closely resemble specimens of *Gryphaea mexicana* Felix illustrated by Felix (1891, pl. 27, figs. 30–30a), Cragin (1905, pl. 3, figs. 1–6), and Imlay (1940, pl. 54, figs. 1–7). As Imlay (1940) stressed, left valves of *G. mexicana* are highly variable, and the specimens from the Little Hatchet Mountains fall within this range of variation. Therefore, but in view of the poor preservation of the specimens from the Little Hatchet Mountains, we tentatively assign them to that taxon as cf. *Gryphaea mexicana*. *Gryphaea mexicana* is known from Upper Jurassic strata in Oaxaca and Durango, Mexico, and the Malone Mountains of west Texas (Felix, 1891; Cragin, 1905; Imlay, 1940).

Gastropods

NMMNH P-31554, collected by P. Harrigan at locality 4470, in a grainstone bed 5 m (16 ft) stratigraphically below P-31555 (Harrigan, 1995), includes parts of three high-spined, slender gastropods whose shells have been partially eroded away to reveal their inner morphology (Fig. 3I–J). The specimens are thin-shelled with slender, solid columella. The aperture, growth lines, and sutural characteristics are not visible due to recrystallization and erosion. One specimen (Fig. 3I) is 40 mm (1.6 inches) long and has an apical angle of 9°. The second specimen (Fig. 3J) is missing the apical angle but is even narrower than the first specimen and has an incomplete length of 44 mm (1.8 inches). The whorl sides are slightly concave to flat. A prominent palatal fold halfway along the whorl sides extends almost halfway

into the whorl. A less prominent, thinner parietal fold occurs near the top of each whorl along the columella. The lower whorls are nearly twice as high as wide.

We identify these specimens as nerineid gastropods due to the presence of the palatal and parietal folds (cf. Cragin, 1905, pp. 96–98, pl. 20, figs. 1–3, pl. 21, figs. 4–5), which are absent in turrnellid gastropods. In addition, turrnellid gastropods have convex whorls that are usually wider than they are high rather than much higher than wide as in the Broken Jug specimens. A more precise identification is not possible, but it is significant that the gastropods from the Little Hatchet Mountains are not turrnellids, as this gastropod family occurs in Cretaceous and younger rocks (Tracey et al., 1993). In contrast, the Nerineidae have a Jurassic–Cretaceous temporal range (Tracey et al., 1993), so they are not precise age indicators at the family level but do not preclude a Jurassic age.

Coral

We identify a coral from locality 4470, NMMNH P-31555 (Fig. 3K–L), as *Thamnasteria* sp. cf. *T. imlayi* Wells, 1942. The corallum of the single specimen is colonial, thamnasteroid, and ramose. The growth form is bushy to phaceloid in appearance. Branches are cylindrical and relatively narrow, 3–7 mm (0.1–0.3 inch) in diameter. Branch bifurcations are frequent, dichotomous, and irregularly spaced. Newly formed branches initially diverge from each other at an angle of ~30°–45° but, with further growth branches from the same stem, often exhibit a general trend toward subparallel orientation. The corallites are numerous, relatively small, closely spaced, and arranged perpendicular to the long axis of the branch. The calices are exclusively monocentric, and the calicular margins are generally circular in outline. Calice diameter ranges from 1.0 to 1.4 mm (0.04–0.06 inch), and the distance between the centers of calices ranges from about 1.4 to 1.8 mm (0.06–0.08 inch). The septa are well developed, solid, about 20–24 in number, and hexamerally arranged in three cycles, the third cycle possibly incomplete. The septa of the first and second cycles are subequal in size and extended nearly to the corallite axis. The septa of the third cycle are weakly developed, very much shorter than previous septa, and possibly incomplete. All septa are continuous beyond the calicular margin as septocostae. The septocostae are then confluent between adjacent corallites. The columella is styliform, deep in the calice, and rarely observed due to recrystallization. Wall or other thecal structures were not observed.

NMMNH P-31555, as preserved, is about 75 mm (3 inches) in height and 160 x 45 mm (6.2 x 1.8 inches) in width. The sub-

parallel alignment of branches on one side of the sample indicates that the actual height of the corallum was at least 120 mm (4.8 inches). Overall, the morphological details of the corallum are poorly preserved. Faint protuberances on the surfaces of some of the branches are relics of the corallites. The septal arrangement was observed only in a few of the exposed corallites on one small part of a branch. Details of the columella were not determined. Skeletal microstructure and internal skeletal architecture are not preserved.

The morphology of NMMNH P-31555 is, with minor exception, the same as that described for *Thamnasteria imlayi* Wells (1942, pp. 127–128, pl. 21, figs. 1–3). *Thamnasteria imlayi* Wells was originally described from samples retrieved from the Jurassic (probably Oxfordian) Smackover Limestone in the subsurface of Columbia

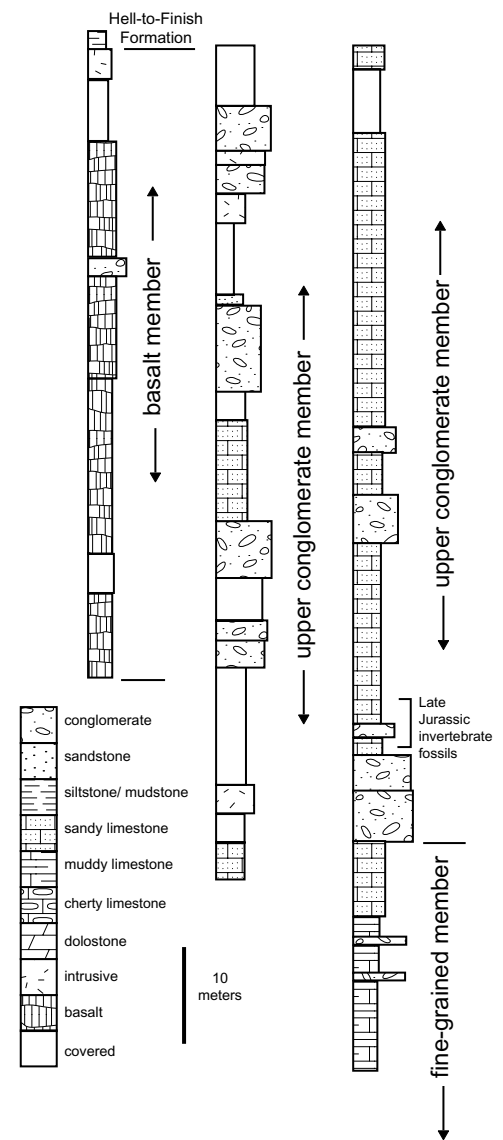
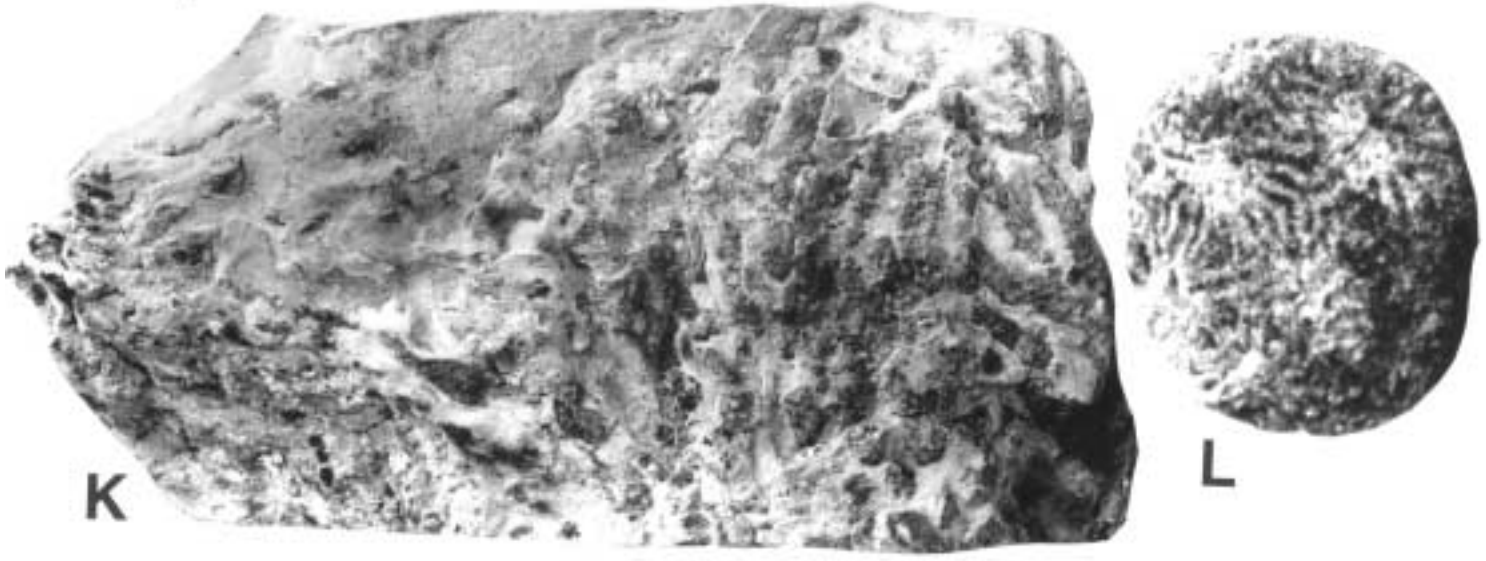
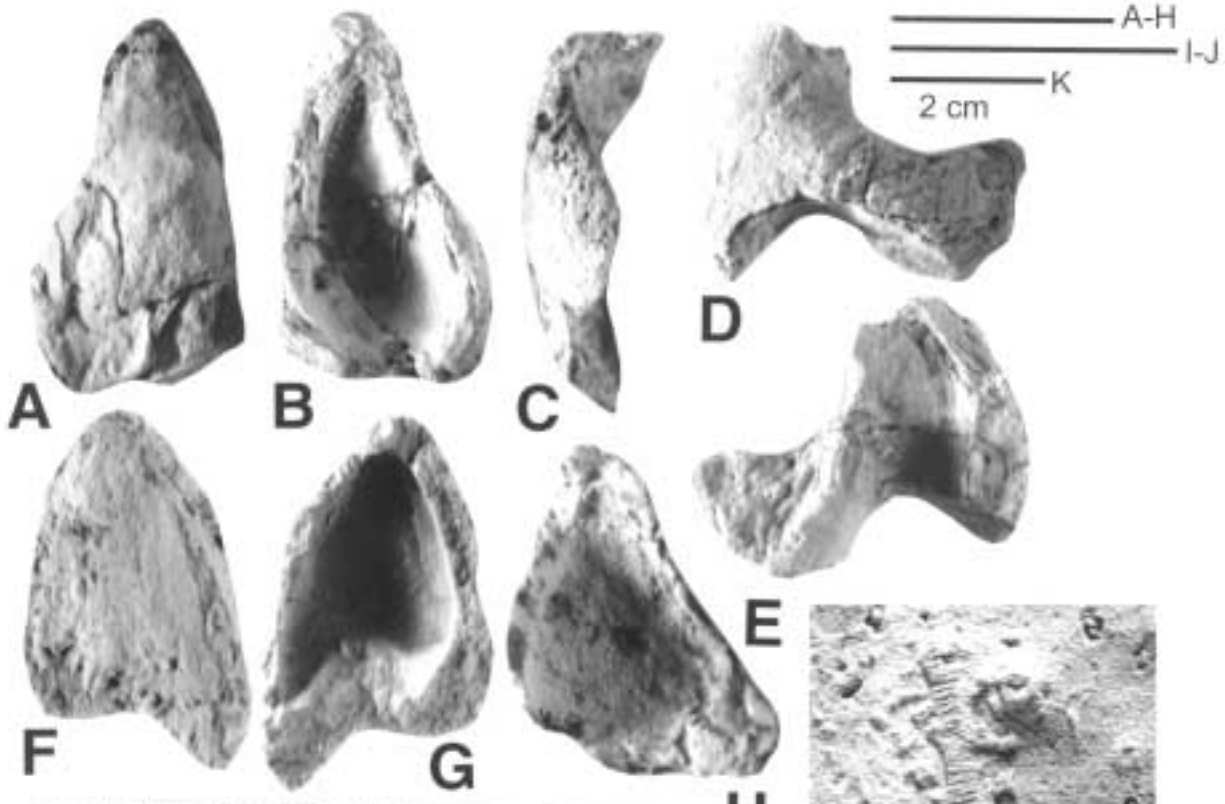


FIGURE 2—Stratigraphic section (right column lowest, left column highest) of part of the Broken Jug Formation showing stratigraphic position of invertebrate fossils described here. Modified from Lucas and Lawton (2000).



County, Arkansas. The branches of the type material are slightly larger in diameter, from 10 to 20 mm (0.4–0.8 inch), but at the distal ends the diameter is only 5 mm (0.2 inch), which is within the range of measurements of the specimen from New Mexico. The diameters of the calices and spacing of the calcular centers of the New Mexico specimen essentially are the same as those of *T. imlayi*. The septal arrangement also appears to be the same. Comparisons with other Late Jurassic and Early Cretaceous members of this genus were given by Wells (1942).

The specimen of *Thamnasteria* sp. cf. *T. lamourouxii* Lesauvage, described by Reyeros de Castillo (1974, p. 16, pl. 3, figs. 1–3) from the Upper Jurassic of the Sierra del Águila, northern Chihuahua, appears to be similar both to the type material of *T. imlayi* and to the New Mexico specimen. The Chihuahua specimen has branches from 6 to 15 mm (0.2–0.6 inch) in diameter, calices 1.5 mm (0.06 inch) in diameter, and 18 septa. It is also interesting to note that the style of preservation of the Chihuahua specimen (see Reyeros de Castillo, 1974, pl. 3, figs. 1, 2) appears to be similar to that of the New Mexico specimen: both coralla are badly recrystallized and internally filled by gray or white sparry calcite. In addition, the sedimentary matrix of both specimens looks very similar.

Discussion

The fossils described here differ fundamentally from the Lower Cretaceous marine invertebrate fossils found in the Aptian–Albian U-Bar Formation, which until now yielded the oldest Mesozoic marine invertebrate fossils found in southwestern New Mexico. U-Bar invertebrate fossil assemblages are diverse, prolific, and dominated by bivalves such as *Exogyra*, *Ostrea*, *Ceratostrongylo*, and trioniids, by rudistids, by turrillid gastropods, and, in some intervals, by ammonites and spatangoid echinoids (e.g., Lucas, 2000a, b; Lucas and Estep, 2000; Lucas and Lawton, 2000). In contrast, the Broken Jug Formation fossils from the Little Hatchet Mountains are of low diversity and abundance and are nerineid gastropods, which range in age

FIGURE 3—Late Jurassic invertebrate fossils from the upper conglomerate member of the Broken Jug Formation, Little Hatchet Mountains. A–H, cf. *Gryphaea mexicana*; A–B, NMMNH P-32959, left valve, external (A) and internal (B) views; C, H, NMMNH P-32958, left valve, side (C) and internal (H) views; D–E, NMMNH P-32957, fragmentary left valve, external (D) and internal (E) views; F–G, NMMNH P-32960, left valve, external (F) and internal (G) views. I–J, Nerineidae, NMMNH P-31554. K–L, *Thamnasteria* sp. cf. *T. imlayi*, NMMNH P-31555, lateral view of corallum (K) and detail of calices (L). Note that L is 12 times larger than K.

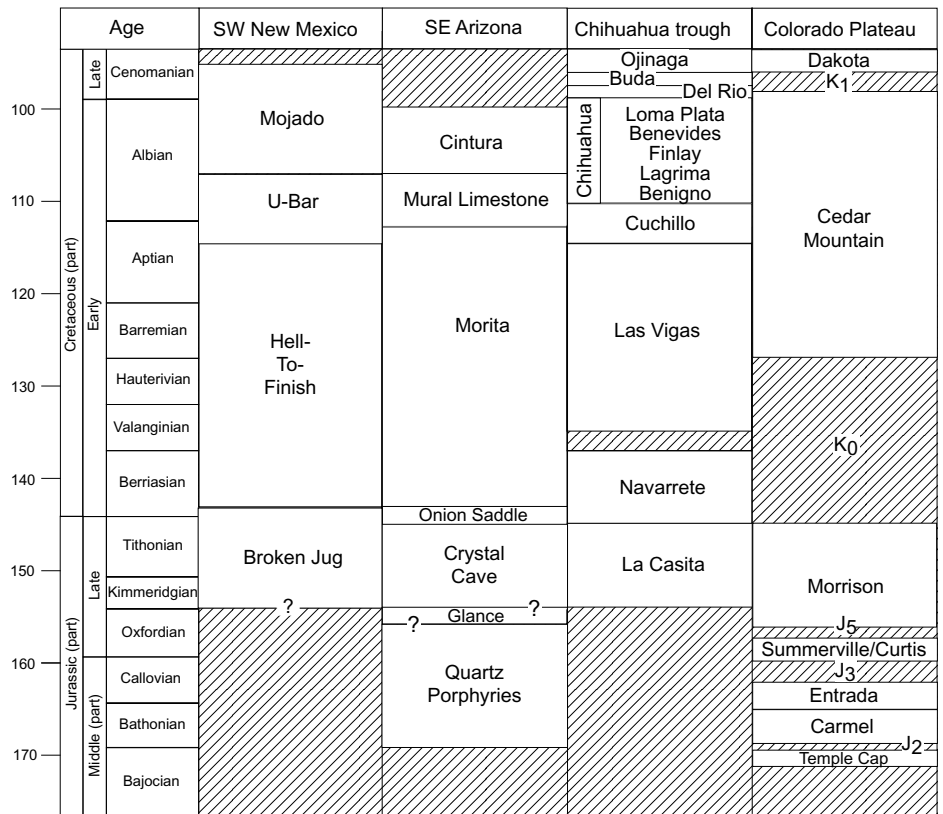


FIGURE 4—Preliminary correlation of Upper Jurassic and Lower Cretaceous stratigraphy of the Bisbee Basin in southwestern New Mexico and southeastern Arizona with strata of the Chihuahua trough (Monreal and Longoria, 1999) and the Colorado Plateau (Kowallis et al., in press).

from Jurassic to Cretaceous, and gryphaeid bivalves and a coral that clearly indicate a Late Jurassic age. They suggest broad correlation to the La Casita Formation of northern Mexico, which, based on its ammonoids, is Kimmeridgian to Tithonian in age (e.g., Monreal and Longoria, 1999).

This age range can be applied to the fossils described here and thus to part of the Broken Jug Formation. The entire Broken Jug Formation has been interpreted to represent a single lithosome deposited near the thick keel of the Bisbee rift basin (Lawton, 2000; Lucas and Lawton, 2000). So, a Late Jurassic age may be reasonable for the entire formation (Fig. 4).

The presence of Late Jurassic marine rocks in southwestern New Mexico (and in southeastern Arizona: Lawton and Olmstead, 1995; Olmstead and Young, 2000) has important implications for interpreting the Late Jurassic paleogeography of the Southwest. The nonmarine Morrison Formation was deposited across the southern Colorado Plateau during Late Jurassic time, and the southern edge of its basin of deposition was long considered to be an east-southeast-trending uplift, the “Mogollon highlands,” in west-central New Mexico and east-central Arizona (Smith, 1951; Harshbarger et al., 1957; Lucas and Anderson, 1997). However, some workers (notably Bilodeau, 1986) have questioned the location of these highlands, placing

them farther south, in northern Mexico and southwestern Arizona. Nevertheless, simultaneous Late Jurassic marine deposition in the Bisbee Basin of southwestern New Mexico-southeastern Arizona and fluvial deposition (with some northerly paleoflow indicators) of the Morrison Formation on the southern Colorado Plateau necessitates a highland between the two depositional systems (Fig. 5).

We regard the Burro uplift (Elston, 1958), lying directly north of the Bisbee Basin, as the drainage divide between north-draining Morrison fluvial systems and contemporary marine deposits to the south. This topographically elevated rift shoulder lay north of the Bisbee Basin in the Early Cretaceous and provided a source for detritus of the Hell-to-Finish Formation (Bilodeau and Lindberg, 1983; Mack et al., 1986), which overlies the Broken Jug Formation (Fig. 4). We infer that the rift shoulder, with a steep southern escarpment facing the Bisbee Basin and a gentle northern flank forming the headwaters of Morrison rivers, was established at least as early as Late Jurassic time.

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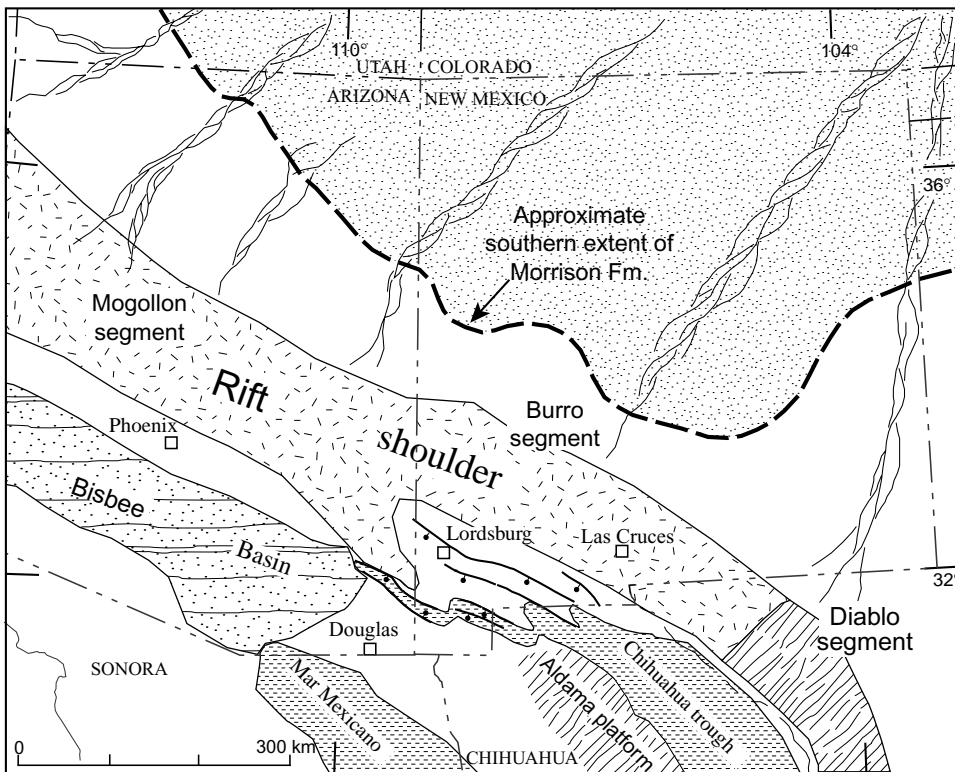


FIGURE 5—Simplified paleogeographic map of Arizona–New Mexico during the Late Jurassic (Kimmeridgian) deposition of the Broken Jug Formation and the Salt Wash Member of the Morrison Formation. Braided fluvial systems drained northeastward from a rift shoulder that lay northeast of the marine embayments of the Chihuahua trough and the Mar Mexicano, which was initiated slightly earlier, in late Oxfordian time, upon cessation of a Triassic–late Middle Jurassic magmatic arc. These two marine basins were separated by the Aldama platform, underlain by upper Paleozoic sedimentary rocks. The rift shoulder is here divided into three segments, the Mogollon, Burro, and Diablo segments. Basal strata of the part of the Bisbee Basin indicated by the sandstone pattern are poorly dated, and their Jurassic distribution is inferred from the extent of Lower Cretaceous rocks of the Bisbee Basin (Bilodeau and Lindberg, 1983). Southern extent of Morrison exposures is an erosional limit beneath the Cenomanian Dakota Sandstone. After Blakey (1989), Peterson (1994), Lucas and Anderson (1997), Robinson and McCabe (1997), Cather (1999), and Lawton (2000).

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